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AMENDMENTS TO THE SPECIFICATION

Please make the following amendments to the specification.

Please replace paragraph [0013] with the following amended paragraph:

[0013] A simple system is used to describe fragments, analogues, and derivatives of GLP-2. For example, Lys²⁰GLP-2(1-33) designates a fragment of GLP-2 formally derived from GLP-2 by deleting the amino acid residues No. 34 and substituting the naturally occurring amino acid residue in position 20 (Arg) by Lys. Similarly, Arg³⁰Lys³⁵(Nε-tetradecanoyl)GLP-1GLP-2(1-35) designates a derivative of a GLP-2 analogue formally derived from GLP-2 by C-terminal addition of a Lys residue, exchange of the naturally occurring amino acid residue in position 30 (Lys) with an Arg residue and tetradecanoylation of the ε-amino group of the Lys residue in position 35.

Please replace paragraph [0016] with the following amended paragraph:

[0016] In a preferred embodiment, the present invention relates to a GLP-2 derivative wherein the parent peptide has an amino acid sequence according to the formula ~~the following amino acid sequence (SEQ ID NO:1):~~
~~X¹ H His Xaa² X² D Asp G Gly S Ser F Phe S Ser D Asp E Glu M Met N Asn F~~
~~Thr Xaa³ X³ L Leu D Asp X⁴ Xaa⁴ L Leu A Ala X⁵ Xaa⁵ X⁶ Xaa⁶ D Asp F Phe I~~
~~Ile N Asn W Trp L Leu X⁷ Xaa⁷ X⁸ Xaa⁸ F Thr K Lys I Ile F Thr D Asp X⁹ Xaa⁹~~
~~Xaa¹⁰ (SEQ ID NO:1),~~

wherein

~~X¹ is NH₂, DFPEEVAIVEELGRR (SEQ ID NO:2), DFPEEVTIVEELGRR (SEQ ID NO:3), DFPEEVNIVEELRRR (SEQ ID NO:4), or a fragment thereof,~~

~~Xaa² X² is Ala or Gly,~~

~~Xaa³ X³ is Ile or Val,~~

~~Xaa⁴ X⁴ is Asn, Ser or His,~~

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Xaa⁵ X⁵ is Ala or Thr,
Xaa⁶ X⁶ is Arg or Lys,
Xaa⁷ X⁷ is Ile or Leu,
Xaa⁸ X⁸ is Gln or His, and
Xaa⁹ X⁹ is OH, Lys, or Arg, and
Xaa¹⁰ is Arg-Lys, Lys-Arg, Arg-Arg or is missing Lys-Lys (Formula I).

In one aspect, the amino acid sequence of the GLP-2 derivative further includes a sequence selected from Asp Phe Pro Glu Glu Val Ala Ile Val Glu Glu Leu Gly Arg Arg (SEQ ID NO:2), Asp Phe Pro Glu Glu Val Thr Ile Val Glu Glu Leu Gly Arg Arg (SEQ ID NO:3), Asp Phe Pro Glu Glu Val Asn Ile Val Glu Glu Leu Arg Arg Arg (SEQ ID NO:4), or a fragment thereof, positioned at the N-terminus of the Formula I sequence (Formula II).

Please replace paragraph [0025] with the following amended paragraph:

[0025] In a particular aspect, the invention relates to use of a pharmaceutical composition comprising a peptide with an the following amino acid sequence according to Formula I or Formula II X²-H-X²-D-G-S-F-S D-E-M-N-T-X³-L-D-X⁴-L-A-X⁵-X⁶-D-F-I-N-W-L-X⁷-X⁸-T-K-I-T-D-X⁹ (SEQ ID NO:1) wherein X² is NH₂, DFPEEVAIVEELGRR (SEQ ID NO:2), DFPEEVIVEELGRR (SEQ ID NO:3), DFPEEVNIVEELRRR (SEQ ID NO:4), or a fragment thereof, X² is Ala or Gly, X³ is Ile or Val, X⁴ is Asn, Ser or His, X⁵ is Ala or Thr, X⁶ is Arg or Lys, X⁷ is Ile or Leu, X⁸ is Gln or His, or X⁹ is OH, Lys, Arg, Arg-Lys, Lys-Arg, Arg-Arg or Lys-Lys for the prophylaxis or treatment of diseases or disorders associated with Impaired appetite regulation.

Please replace "aspect 47" (on page 24 of the specification) with the following amended aspect:

47. A pharmaceutical composition of any of aspects 37-46, wherein the parent peptide has an the following amino acid sequence according to

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Formula I or Formula II (SEQ ID NO:1)

~~X¹-H-X²-D-G-S-F-S-D-E-M-N-T-X³-L-D-X⁴-L-A-X⁵-X⁶-D-F-I-N-W-L-X⁷-X⁸-T-K-I-T-D-X⁹~~

wherein

~~X¹ is NH₂, DFPEEVAIVEELGRR (SEQ ID NO:2), DFPEEVITIVEELGRR (SEQ ID NO:3), DFPEEVNIVEELRRR (SEQ ID NO:4), or a fragment thereof,~~

~~X² is Ala or Gly,~~

~~X³ is Ile or Val,~~

~~X⁴ is Asn, Ser or His,~~

~~X⁵ is Ala or Thr,~~

~~X⁶ is Arg or Lys,~~

~~X⁷ is Ile or Leu,~~

~~X⁸ is Gln or His, and~~

~~X⁹ is OH, Lys, Arg, Arg-Lys, Lys-Arg, Arg-Arg or Lys-Lys.~~

Please replace aspects "58-91" on pages 25-27 of the application with the following substitute aspects:

58. Use of a pharmaceutical composition comprising a peptide with the following an amino acid sequence according to Formula I or Formula II ~~X¹-H-X²-D-G-S-F-S-D-E-M-N-T-X³-L-D-X⁴-L-A-X⁵-X⁶-D-F-I-N-W-L-X⁷-X⁸-T-K-I-T-D-X⁹~~ wherein X¹ is NH₂, DFPEEVAIVEELGRR, DFPEEVITIVEELGRR, DFPEEVNIVEELRRR, or a fragment thereof, X² is Ala or Gly, X³ is Ile or Val, X⁴ is Asn, Ser or His, X⁵ is Ala or Thr, X⁶ is Arg or Lys, X⁷ is Ile or Leu, X⁸ is Gln or His, and X⁹ is OH, Lys, Arg, Arg-Lys, Lys-Arg, Arg-Arg or Lys-Lys together with a pharmaceutically acceptable excipient or vehicle for appetite suppression or satiety ~~satiety~~ induction.

~~59-69.~~ The use of a composition according to aspect ~~58~~ 59, wherein the amino acid sequence is according to Formula I X¹ is NH₂.

~~60-70.~~ The use of a composition according to aspect 59, wherein X² Xaa² is Ala.

~~61-71.~~ The use of a composition according to aspect 59, wherein X³ Xaa³ is Ile.

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- ~~62~~ ~~72~~. The use of a composition according to aspect 59, wherein
~~X4~~ Xaa⁴ is Asn.
- ~~63~~ ~~73~~. The use of a composition according to aspect 59, wherein
Xaa⁵ ~~X5~~ is Ala.
- ~~64~~ ~~74~~. The use of a composition according to aspect 59, wherein
Xaa⁶ ~~X6~~ is Arg.
- ~~65~~ ~~75~~. The use of a composition according to aspect 59, wherein
~~X7~~ Xaa⁷ is Ile.
- ~~66~~ ~~76~~. The use of a composition according to aspect 59, wherein
Xaa⁸ ~~X8~~ is Gln.
- ~~67~~ ~~77~~. The use of a composition according to aspect 59, wherein
~~X9~~ Xaa⁹ is OH.
- ~~68~~ ~~78~~. The use of a composition according to aspect 59, wherein
the peptide has the sequence
HADGSFSDEMNTILDNLAA~~R~~DFIQTKITD (SEQ ID NO:5),
HADGSFSDEMNTILDNLAT~~R~~DFINWLIQTKITD (SEQ ID NO:6), or
HADGSFSDEMNTVLDNLAT~~R~~DFINWLLHTKITD (SEQ ID NO:7).
- ~~69~~ ~~79~~. The use of a composition according to any of aspects 59-
~~68~~ ~~71~~, for the prophylaxis or treatment of diseases or disorders associated
with impaired appetite regulation.
- ~~70~~ ~~80~~. The use of a composition according to any of the aspects
59-~~69~~ ~~70~~ for the prophylaxis or treatment of obesity or type II diabetes.
- ~~71~~ ~~81~~. A pharmaceutical composition comprising a peptide of any
of the compositions used in any of aspects 59-70 in combination with
another appetite-suppressing or satiety-inducing agent.
- ~~72~~ ~~82~~. A composition according to aspect ~~71~~ ~~73~~, wherein said
other appetite suppressing or satiety-inducing agent is glucagon-like
peptide-1.
- ~~73~~ ~~83~~. A method of treating diseases or disorders associated with
impaired appetite regulation, the method comprising administering to an
individual in need of such treatment an amount of a peptide comprised in
any of the compositions used according to any of aspects 59-70 sufficient to
suppress appetite or induce satiety in said individual.

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74 84. A method according to aspect 73 83, wherein the disease or disorder is obesity or type II diabetes.

75 85. A method according to aspect 73 83, wherein the amount of the peptide is in the range of from about 10 μ g/kg body weight to about 5 mg/kg body weight.

76 86. A method of treating diseases or disorders associated with impaired appetite regulation, the method comprising administering to an individual in need of such treatment an amount of a peptide comprised in the composition used according to aspect 59 sufficient to suppress appetite or induce satiety in said individual.

77 87. A method according to aspect 76 86, wherein the disease or disorder is obesity or type II diabetes.

78 88. A method according to aspect 76 86, wherein the amount of the peptide is in the range of from about 10pg/kg body weight to about 5 mg/kg body weight.

~~89. A method of treating diseases or disorders associated with impaired appetite regulation, the method comprising administering to an individual in need of such treatment an amount of a fraction according to aspect 86 sufficient to suppress appetite or induce satiety in said individual.~~

~~90. A method according to aspect 89, wherein the disease or disorder is obesity or type II diabetes.~~

79 91. Use of a peptide comprised in a composition used according to any of aspects 59-70 for the manufacture of a medicament for the prophylaxis or treatment of diseases or disorders associated with impaired appetite regulation.